AMENDMENTS TO THE CLAIMS

This listing of the claims replaces all prior versions and listings:

- 1. (original): A composition comprising a substantially integral bARE class protein.
- 2. (original): The composition according to claim 1, wherein the composition comprises a stabilising agent.
- 3. (original): The composition according to claim 2, wherein the stabilising agent is a charged amino acid or an analogue thereof.
- 4. (currently amended): The composition according to claim 3, wherein the stabilising agent is Arginine or Arginine Phosphate.
- 5. (original): The composition according to claim 4, wherein the Arginine or Arginine phosphate is present in an amount of from about 100mM to about 400mM.
- 6. (original): The composition according to claim 2, wherein the composition comprises an uncharged agent or an analogue thereof.
- 7. (original): The composition according to claim 6, wherein the composition comprises a zwitterionic agent.
- 8. (original): The composition according to claim 7, wherein the zwitterionic agent is a zwitterionic detergent.
- 9. (original): The composition according to claim 8, wherein the zwitterionic detergent is 3-(3-Cholamidopropyl)-dimethylammonio-1-propanesulfonate (CHAPS).
- 10. (original): The composition according to claim 9, wherein the zwitterionic detergent is present in an amount of from about 0.05% to about 0.5% by weight per volume (w/v).

- 11. (currently amended): The composition according to claim 2, wherein the composition comprises a charged amino acid or analogue according to any one of claims 3-5 and an uncharged agent according to any one of claims 6-9.
- 12. (currently amended): The composition according to claim <u>1</u> any one of claims <u>1</u>-11, wherein the Integrity of the bARE class protein is determined with reference to an Integrity Ratio.
- 13. (currently amended): The composition according to <u>claim 1</u> any one of claims 1-12, wherein the bARE protein is an AB5 protein.
- 14. (original): The composition according to claim 13, wherein the bARE protein is an LTK63 or LTK 72 protein.
- 15. (original): A method of stabilising a bARE protein wherein the method comprises providing a bARE class protein and combining the bARE class protein with a stabilising agent.
- 16. (original): The method according to claim 15, wherein the stabilising agent is a charged amino acid or an analogue thereof.
- 17. (original): The method according to claim 16, wherein the stabilising agent is Arginine or Arginine Phosphate.
- 18. (original): The method according to claim 17, wherein the Arginine or Arginine phosphate is present in an amount of from about 100mM to about 400mM.
- 19. (original): The method according to claim 15, wherein the stabilising agent is an uncharged agent.
- 20. (original): The method according to claim 19, wherein the uncharged agent is a zwitterionic agent.

- 21. (original): The method according to claim 20, wherein the zwitterionic agent is a zwitterionic detergent.
- 22. (original): The method according to claim 21, wherein the zwitterionic detergent is 3-(3-Cholamidopropyl)-dimethylammonio-1-propanesulfonate (CHAPS).
- 23. (original): The method according to claim 22, wherein the zwitterionic detergent is present in an amount of from about 0.05% to about 0.5% by weight per volume (w/v).
- 24. (currently amended): The method according to claim 15, wherein the stabilising agent comprises a charged amino acid according to any one of claims 16-18 and an uncharged agent-according to any one of claims 19-23.
- 25. (currently amended): The method according to <u>claim 15</u> any one of claims 15-24, wherein the stabilising of the bARE class protein is determined with reference to an Integrity Ratio.
- 26. (currently amended): The method according to <u>claim 15</u> any one of claims 15-25, wherein the bARE protein is an AB5 protein.
- 27. (original): The method according to claim 26, wherein the AB5 protein is an LTK63 or LTK 72 protein.
- 28. (currently amended): A method of analysing a bARE class protein <u>comprising</u> analysing a composition comprising the bARE class protein under non-dissociating conditions to which differentiate between integral and dissociated bARE class proteins.
- 29. (currently amended): The method according to claim 28, wherein the method comprises a separation step on separating the proteins using a charged polymeric separation material.

- 30. (original): The method according to claim 29, wherein the polymeric separation material is a hydrogel monomer.
- 31. (original): The method according to claim 30, wherein the hydrogel monomer is a hydroxylated polymethacrylate (HEMA) monomer.
- 32. (original): The method according to claim 31, wherein the HEMA has a particle size of about 6 microns.
- 33. (currently amended): The method according to claim 31-or 32, wherein the HEMA has a porosity of about 250A.
- 34. (original): A method of analysing a bARE class protein wherein the method comprises:
- (i) applying a bARE class protein to a charged polymeric separation material in an apparatus configured to resolve an integral bARE class protein from a dissociated bARE class protein;
- (ii) treating the separation material comprising the applied bARE class protein with an ionic buffer; and
 - (iii) detecting one or more integral or dissociated bARE class proteins.
- 35. (currently amended): The method according to claim 34, wherein the separation material is a hydrogel monomer as defined in any one of claims 30-33.
- 36. (currently amended): The method according to claim 34 or 35, wherein the ionic buffer is a physiologically acceptable buffer with a pH of from about 7.0 to about 8.0.
- 37. (original): A method for identifying a bARE class protein stabilisation agent wherein the method comprises:
- (i) combining a bARE class protein with a candidate stabilising agent to form a bARE protein sample;

- (ii) applying the bARE protein sample to a charged polymeric separation material in an apparatus configured to resolve an integral bARE class protein from a dissociated bARE class protein;
- (iii) treating the separation material comprising the applied bARE class protein with an ionic buffer;
 - (iv) detecting one or more integral or dissociated bARE class proteins; and
- (v) determining whether the candidate stabilising agent is a bARE protein stabilising agent.
- 38. (original): The method according to claim 37, wherein the method comprises calculating an Integrity Ratio for the bARE protein sample.
- 39. (original): The method according to claim 38, wherein the method further comprises comparing the Integrity Ratio for the bARE protein sample with an Integrity Ratio for a control without a candidate stabilising agent.
- 40. (currently amended): A stabilising agent identified by the method of claim 37 any one of claims 37-39.
- 41. (original): The stabilising agent according to claim 40, which is a functional stabilising agent.
- 42. (original): The stabilising agent according to claim 40, which is a physical stabilising agent.
- 43. (currently amended): An immunogenic composition comprising a composition according to claim 1-any one of claims 1-14.
- 44. (original): An immunogenic composition according to claim 43, wherein further comprising an adjuvant, wherein said adjuvant is not the bARE protein.
- 45. (original): An immogenic composition according to claim 44, wherein the adjuvant is a mucosal adjuvant.

- 46. (canceled).
- 47. (currently amended): A method of treating a mammal to prevent and/or treat an immune disorder comprising administering a composition according to <u>claim 43</u> any one of <u>claims 43-45</u>.
 - 48. (original): A method according to claim 47 wherein the mammal is a human.
 - 49 to 60. (canceled).